Amendments to the Claims:

The following listing of claims will replace all prior versions, and listings, of claims in the application:

1. (Original) A compound of formula (I)

$$X \xrightarrow{N} \xrightarrow{Q} \xrightarrow{Q} \xrightarrow{N} \xrightarrow{N} \xrightarrow{R} \xrightarrow{E}_{m} \xrightarrow{B} \xrightarrow{D}_{h}$$

$$(I)$$

or a pharmaceutically acceptable salt or ester thereof, wherein

X is

- 1) H,
- 2) aryl,
- 3) heteroaryl or
- 4) a group of formula

wherein aryl and heteroaryl can be unsubstituted or substituted with 1 to 4 substituents selected from \mathbb{R}^{a} , as defined hereinafter;

Y is

- 1) H,
- 2) (C_1-C_6) alkyl,
- 3) (C₃-C₇)cycloalkyl or
- 4) (C_3-C_7) cycloalkyl- (C_1-C_3) alkyl;

Q is

- 1) aryl,
- 2) aryl- (C_1-C_6) alkyl,
- 3) heteroaryl or
- 4) heteroaryl- (C_1-C_6) alkyl;

wherein aryl and heteroaryl can be optionally substituted with 1 to 3 substituents selected from \mathbb{R}^{a} ; and alkyl can be optionally substituted with Cy;

Cy is cycloalkyl, heterocyclyl, aryl or heteroaryl;

A is

1) (C_1-C_6) alkyl,

- 2) (C₂-C₆)alkenyl,
- 3) (C₂-C₆)alkynyl,
- 4) Cy or
- 5) Cy- (C_1-C_6) alkyl;

wherein alkyl and cycloalkyl can be optionally substituted with 1 to 2 substituents selected from \mathbf{R}^c , as defined hereinafter; and Cy can be optionally substituted with 1 to 3 substituents selected from \mathbf{R}^a ;

B is

- 1) N or
- 2) C(D);

D is independently

- 1) H,
- 2) halogen,
- 3) (C_1-C_6) alkyl,
- 4) (C_2-C_6) alkenyl,
- 5) (C₂-C₆)alkynyl,
- 6) $-NR^bR^b$,
- 7) $-NO_2$ or
- 8) -CN;

wherein R^b is to be defined hereinafter;

E is

- 1) CH₂,
- 2) CHR^b or
- 3) CR^bR^c;

R1 is

- 1) H,
- 2) (C_1-C_6) alkyl,
- 3) (C_2-C_6) alkenyl,
- 4) (C₂-C₆)alkynyl,
- 5) Cy,
- 6) Cy- (C_1-C_3) alkyl,
- 7) $-(CH_2)_kC(O)NR^bR^b$ or
- 8) (C_1-C_6) alkoxy (C_1-C_6) alkyl;

wherein Cy can be unsubstituted or substituted with a group selected from $\mathbf{R}^{\mathbf{a}}$ and alkyl, alkynyl and alkoxy can be unsubstituted or substituted with a group selected from $\mathbf{R}^{\mathbf{c}}$;

R2 is

- 1) H,
- 2) (C_1-C_9) alkyl,
- 3) (C₂-C₉)alkenyl,
- 4) (C₂-C₉)alkynyl,

- 5) Cy or
- 6) Cy-(C₁-C₃)alkyl;

wherein Cy can be unsubstituted or substituted with a group selected from R^a and alkyl, alkenyl and alkynyl can be unsubstituted or substituted with a group selected from R^c ;

R3 is

- 1) H or
- 2) (C_1-C_6) alkyl;

R^a is independently

- 1) H,
- 2) halogen,
- 3) (C_1-C_6) alkyl,
- 4) (C_2-C_6) alkenyl,
- 5) (C_2-C_6) alkynyl,
- 6) Cy,
- 7) -ORb,
- $8) -SR^b$
- 9) $-NR^bR^b$,
- 10) $-NR^bC(N)NR^bR^b$.
- 11) $-C(O)R^{b}$,
- 12) $-C(O)NR^bR^b$.
- 13) $-NC(O)R^{b}$,
- 14) $-SO_2NR^bR^b$,
- 15) $-NO_2$,
- 16) -CN,
- 17) $-CF_3$ or
- 18) amino- (C_1-C_6) alkyl;

R^b is independently

- 1) H,
- 2) (C_1-C_6) alkyl,
- 3) (C_2-C_6) alkenyl,
- 4) (C₂-C₆)alkynyl,
- 5) (C₃-C₇)cycloalkyl,
- 6) aryl,
- 7) heteroaryl,

or in the context of D, R1, R^a and R^c, R^b and R^b together with the atom to which they are attached can also form a 5 to 6 membered ring containing 1 to 2 heteroatoms selected from N, O and S;

R^c is independently

- 1) H,
- 2) halogen,
- 3) Cy,

- 4) –CN,
- $-OR^{b}$
- $-SR^b$
- 7) $-NR^bR^b$ or
- 8) $-NR^bC(N)NR^bR^b$;

k is an integer 0 or 1;

h is an integer from 0 to 4;

n is an integer 0 or 1;

m is an integer from 0 to 3;

with the proviso that the compound of formula I is not the compound

and provided that A in formula (I) is not 2-hydroxyethyl.

2. (Original) A compound according to claim 1, wherein

R2 is

- 1) H,
- (C_1-C_6) alkyl,
- (C_2-C_6) alkenyl,
- 4) (C_2-C_6) alkynyl,
- 5) Cy or
- 6) $\text{Cy-}(C_1-C_3)\text{alkyl};$

wherein Cy can be unsubstituted or substituted with a group selected from \mathbb{R}^a and alkyl, alkenyl and alkynyl can be unsubstituted or substituted with a group selected from \mathbb{R}^c .

3. (Currently Amended) A compound according to claim 1-or-2, wherein the compound is a compound of formula IA

or a pharmaceutically acceptable salt or ester thereof,

IA

wherein A, Q, X, Y and n are as defined in claim 1 or claim 2.

4. (Currently Amended) A compound according to claim 1-or-2, wherein the compound is a compound of formula IB

$$X \xrightarrow{N} \begin{array}{c} O \\ N \\ N \\ R \end{array}$$

$$(R) \qquad \qquad (R)$$

 \mathbf{IB}

or a pharmaceutically acceptable salt or ester thereof,

wherein A, D, E, X, Y, h, m and n are as defined in claim 1-or claim 2;

Q is aryl- (C_1) alkyl or heteroaryl- (C_1) alkyl, where aryl or heteroaryl are optionally substituted with 1 to 2 substituents selected from R^a .

5. (Currently Amended) A compound according to claim 1-or-2, wherein the compound is a compound of formula IC

IC

or a pharmaceutically acceptable salt or ester thereof,

wherein R2, A, D, E, Q, h, m and n are as defined in claim 1.

6. (Currently Amended) A compound according to claim 1—or 2, wherein the compound is a compound of formula ID

$$X \xrightarrow{H} Q \xrightarrow{N} M \xrightarrow{N} M \xrightarrow{M} D \xrightarrow{N} M$$

ID

or a pharmaceutically acceptable salt or ester thereof,

wherein A, X, D and h are as defined in claim 1-or-claim 2;

Q is aryl- (C_1) alkyl or heteroaryl- (C_1) alkyl, where aryl or heteroaryl are optionally substituted with 1 to 2 substituents selected from \mathbb{R}^a ; and

m is an integer 1 or 2.

- 7. (Currently Amended) A compound according to claim 1—or 2, wherein the compound of formula I is any of the compounds no 1 to 15 or 23 to 62 as described in the Examples.
- 8. (Currently Amended) A compound according to claim 1-or-2, wherein the compound of formula I is (2R, 2'R)-5-Amino-2-{3'-naphthalen-1-yl-2'-[3-phenethyl-3-(2-pyridin-2-ylethyl)ureido]propionylamino}pentanamide, (2R)-N-(4-Aminobutyl)-3-(1H-indol-3-yl)-2-[3-(3-phenylpropyl)-3-(2-pyridin-2-ylethyl)ureido]propionamide, (2S, 2'R)-2-{2-[3,3-Bis(2-pyridin-2-ylethyl)ureido]-3-naphthalen-1-ylpropionylamino}-4-methylsulfanylbutyramide, (2S, 2'R)-4-Methylsulfanyl-2-{3'-naphthalen-1-yl-2'-[3-phenethyl-3-(2-pyridin-2-ylethyl)ureido]propionylamino}butyramide, (2S, 2'R)-3-Methyl-2-{3'-naphthalen-1-yl-2'-[3-phenethyl-3-(2-pyridin-2-ylethyl)ureido]-propionylamino}butyramide, (2R)-N-cyclohexyl-3-naphthalen-1-yl-2'-[3-phenethyl-3-(2-pyridin-2-ylethyl)ureido]propionamide, (2S,2'R)-2-{3'-naphthalen-1-yl-2'-[3-phenethyl-3-(2-pyridin-2-ylethyl)ureido]propionylamino}-3-phenylpropionamide or (2S,2'R)-2-{2-[3,3-bis(2-pyridin-2-ylethyl)ureido]-3'-naphthalen-1-ylpropionylamino}-3-methylbutyramide.
- 9. (Currently Amended) A compound according to any of the claims 1 to 8 claim 1 where the compound is an SSTR1 selective agonist.
- 10. (Currently Amended) A compound according to any of the claims 1 to 8 claim 1 where the compound is an SSTR1 selective antagonist.
- 11. (Currently Amended) A pharmaceutical composition comprising as active ingredient at least one compound according to any of the claims 1 to 10 claim 1 and at least one pharmaceutically acceptable carrier.

- 12. (Currently Amended) Use of a compound according to any of the claims 1 to 10 for the manufacture of a pharmaceutical preparation A method for the treatment and/or prevention of a disease or condition responding to targeting with a selective SSTR1 compound, comprising administering to a patient with such a disease or condition a compound according to claim 1.
- 13. (Currently Amended) The-use-method according to claim 12, wherein the said disease or condition is a central nervous system disease or disorder, a disease or condition benefiting from the use of anti-proliferative agents, pathological condition in the retina and/or iris-ciliary body, diabetic complication, cancer or excessive proliferation of normal or malignant tissue.
- 14. (Currently Amended) The <u>use</u> method according to claim 12, wherein the said disease or condition is anxiety, depression or schizophrenia.
- 15. (Currently Amended) The <u>use</u> <u>method</u> according to claim 12, wherein the said disease or condition is prostatic cancer, benign prostatic hyperplasia, pancreatic cancer, thyroid cancer, brain tumor or gastro-intestinal tumor.
- 16. (Currently Amended) The—use—method according to claim 12, wherein the said disease or condition is diabetic retinopathy, diabetic nephropathy or diabetic neuropathy.
- 17. (Currently Amended) The <u>use method</u> according to claim 12, wherein the said disease or condition is angiogenesis, vascular restensis, smooth muscle proliferation, endothelial cell proliferation, new blood vessel sprouting or neovascularization.
- 18. (Currently Amended) Use of a compound of according to any of the claims 1-10 in combination with a detectable label, A method for targeting tissues bearing SSTR1s for tissue imaging, comprising administering a compound according to claim 1 in combination with a detectable label.
- 19. (Currently Amended) Use of a compound of according to any of the claims 110 as a carrier for another A method for targeting a therapeutically active compound to be targeted to tissues bearing SSTR1s, comprising administering said therapeutically active compound with a compound according to claim 1 as a carrier for said therapeutically active compound.